

## CLAIMS

1. A prophylactic antimigraine agent comprising as an active ingredient a selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors.

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2. A prophylactic antimigraine agent as claimed in Claim 1, wherein the selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors comprises a) a 5-HT<sub>2B</sub> receptor antagonistic compound as a first ingredient having a selective binding affinity to the 5-HT<sub>2B</sub> receptor, and b) a 5-HT<sub>7</sub> receptor antagonistic compound as a second  
10 ingredient having a selective binding affinity to the 5-HT<sub>7</sub> receptor.

3. A prophylactic antimigraine agent as claimed in Claim 1, wherein the selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors comprises a dual antagonistic compound for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors having a selective binding affinity to both  
15 of the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors.

4. A combined prophylactic preparation for migraine which comprises a) a first pharmaceutical preparation comprising as an active ingredient a 5-HT<sub>2B</sub> receptor antagonistic compound having a selective binding affinity to the 5-HT<sub>2B</sub> receptor, and b) a  
20 second pharmaceutical preparation comprising as an active ingredient a 5-HT<sub>7</sub> receptor antagonistic compound having a selective binding affinity to the 5-HT<sub>7</sub> receptor, and wherein the first and second preparations are administered simultaneously or separately.

5. A prophylactic antimigraine agent as claimed in Claim 1, wherein the  
25 binding affinity for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors is respectively one-hundredth or less to the  $\alpha_1$ , M<sub>1</sub>, D<sub>2</sub>, 5-HT<sub>1A</sub>, 5-HT<sub>1B</sub>, 5-HT<sub>2A</sub>, 5-HT<sub>2C</sub>, 5-HT<sub>3</sub>, 5-HT<sub>4</sub> and 5-HT<sub>6</sub> receptors.

6. Use of the selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors for the manufacture of a prophylactic antimigraine agent.

5 7. Use of "a 5-HT<sub>2B</sub> receptor antagonistic compound having a selective binding affinity to the 5-HT<sub>2B</sub> receptor" for the manufacture of a prophylactic antimigraine agent comprising as an active ingredient a selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors.

10 8. Use of "a 5-HT<sub>7</sub> receptor antagonistic compound having a selective binding affinity to the 5-HT<sub>7</sub> receptor" for the manufacture of a prophylactic antimigraine agent comprising as an active ingredient a selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors.

15 9. A method for prophylaxis of migraine which comprises administering a therapeutically effective amount of a selective dual antagonist for the 5-HT<sub>2B</sub> and 5-HT<sub>7</sub> receptors to a patient.

20 10. A method for prophylaxis of migraine which comprises administering a combination comprising a pharmaceutical preparation containing as an active ingredient a 5-HT<sub>2B</sub> selective receptor antagonistic compound and a pharmaceutical preparation containing as an active ingredient a 5-HT<sub>7</sub> receptor selective antagonistic compound, simultaneously or separately to a patient.